ULTRA SHADE SUNSCREEN SPF 30 TAHERI MD- titanium dioxide, zinc oxide cream LA LASER CENTE, PC, CALIFORNIA PROFESSIONAL MEDICAL CORP

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Ultra Shade Sunsccreen SPF 30

Active Ingredients Purpose

Titanium Dioxide 5.00% Sunscreen

Zinc Oxid 10.00%

Uses

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[Helps prevent sunburn]

Keep out of reach of children. If swallowed get help or contact a Poison Control Center right away.

Stop use and ask a doctor Sif rash occurs

Warnings

Skin Cancer/Skin Aging Alert: Spending time in the sun increases your risk of skin cancer and early skin aging. This product has been shown only to prevent sunburn, not skin cancer or early skin aging.

- **For external use only.**
- Do not use on damage or broken skin.
- When using this product keep out of eyes. Rinse with water to remove

Directions

- ¶Apply generously 15 minutes before sun exposure
- Reapply:
- -after 80 minutes of swimming or sweating
- -immediatley after towel drying
- at least every 2 hours
- Chirldren under 6 months of age: ask a doctor

Acrylates/C10-30 Alkyl Acrylates Crosspolymer, Alumina, Aqua, Butylene Glycol, C12-15 Alkyl Benzoate, C13-14 Isoparaffin, Cetearyl Alcohol,

Coco-Caprylate/Caprate, Coconut Alkanes, Cyclopentasiloxane, Dimethicone, Ethylhexylglycerin, Glycerin, Glyceryl Stearate, Iron Oxides, Laureth-7, Octoyldodecyl Neopentanoate, Polyhydroxystearic Acid, Triethanolamine, Xanthan Gum

TAHERI MD

Ultra Shade Sunscreen SPF 30

Non- Comedogenic

1.70z / 50 ml

TAHERI MD

Ultra Shade Sunscreen

SPF 30

Non-Comedogenic

1.7oz / 50ml

Drug Facts

Active Ingredients

Purpose

Titanium Dioxide 5.00% Zinc Oxide 10.00%

Sunscreen

Uses

· helps prevent sunburn

Warnings

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- · Reapply:
- · after 80 minutes of swimming or sweating
- · immediately after towel drying
- · at least every 2 hours.
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Inactive ingredients

Acrylates/C10-30 Alkyl Acrylates Crosspolymer, Alumina, Aqua, Butylene Glycol, C12-15 Alkyl Benzoate, C13-14 Isoparaffin, Cetearyl Alcohol, Coco-Caprylate/Caprate, Coconut Alkanes, Cyclopentasiloxane, Dimethicone, Ethylhexylglycerin, Glycerin, Glyceryl Stearate, Iron Oxides Laureth-7, Octyldodecyl Neopentanoate, PEG-100 Stearate, Phenoxyethanol, Polyacrylamide, Polyhydroxystearic Acid, Polysorbate 60, Stearic Acid, Triethanolamine, Xanthan Gum

Other information

Protect from excessive heat and direct sunlight.

DISTRIBUTED BY
DANIEL TAHERI, MD
LOS ANGELES, CA 90025
WWW.LALASERCENTER.COM



ULTRA SHADE SUNSCREEN SPF 30 TAHERI MD

titanium dioxide, zinc oxide cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70919-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP) (TITANIUM DIO XIDE - UNII:15FIX9 V2JP)	TITANIUM DIO XIDE	5 g in 100 mL	
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	10 g in 100 mL	

Inactive Ingredients	
Ingredient Name	Strength
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
ALUMINUM O XIDE (UNII: LMI26O6933)	
WATER (UNII: 059QF0KO0R)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
C13-14 ISOPARAFFIN (UNII: E4F12ROE70)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
COCO-CAPRYLATE/CAPRATE (UNII: 8 D9 H4QU9 9 H)	
COCONUT ALKANES (UNII: 1E5KJY107T)	
CYCLOMETHICONE 5 (UNII: 0 THT5PCI0 R)	
DIMETHICO NE (UNII: 92RU3N3Y1O)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
GLYCERIN (UNII: PDC6A3C0OX)	
GLYCERYL MONOSTEARATE (UNII: 230 O U9 XXE4)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
LAURETH-7 (UNII: Z95S6G8201)	
OCTYLDODECYL NEOPENTANOATE (UNII: X8725R883T)	
PEG-100 STEARATE (UNII: YD01N1999R)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POLYACRYLAMIDE (1500 MW) (UNII: 5D6TC4BRWV)	
POLYHYDROXYSTEARIC ACID (2300 MW) (UNII: YXH47AOU0F)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TROLAMINE (UNII: 9O3K93S3TK)	
XANTHAN GUM (UNII: TTV12P4NEE)	

	Packaging			
:	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:70919-001- 01	50 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	08/17/2016	

Marketing Infor	mation		
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date

OTC monograph final	part352	08/17/2016	

Labeler - La Laser cente, pc, california professional medical corp (080323225)

Registrant - LA LASER CENTE, PC, CALIFORNIA PROFESSIONAL MEDICAL CORP (080323225)

Establishment				
Name	Address	ID/FEI	Business Operations	
VEGE-KURL, INC		021072509	manufacture (70919-001)	

Revised: 1/2017

LA LASER CENTE, PC, CALIFORNIA PROFESSIONAL MEDICAL CORP